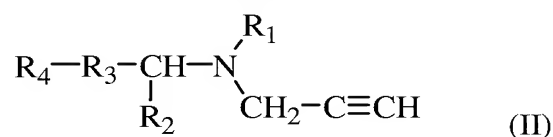


**LISTING OF THE CLAIMS**

This listing of claims will replace all prior versions of the claims and listing of the claims in the application:

1.-13. **(Cancelled)**

14. **(Previously Presented)** A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, wherein the deprenyl compound is represented by the structure:



in which

R<sub>1</sub> is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxy carbonyl, or aryloxy carbonyl;

R<sub>2</sub> is hydrogen or alkyl;

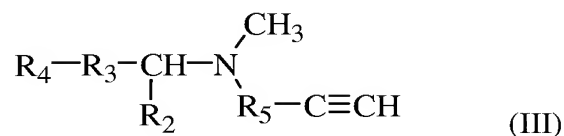
R<sub>3</sub> is a bond or methylene; and

R<sub>4</sub> is aryl or aralkyl; or

R<sub>2</sub> and R<sub>4</sub>-R<sub>3</sub> are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group;

and pharmaceutically acceptable salts thereof.

15. **(Previously Presented)** A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, wherein the deprenyl compound is represented by the structure:



in which

R<sub>2</sub> is hydrogen or alkyl;

R<sub>3</sub> is a bond or methylene; and

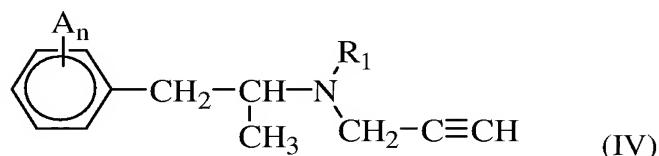
R<sub>4</sub> is aryl or aralkyl; or

R<sub>2</sub> and R<sub>4</sub>-R<sub>3</sub> are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group; and

R<sub>5</sub> is alkylene, alkenylene, alkynylene and alkoxylenes;

and pharmaceutically acceptable salts thereof.

16. **(Previously Presented)** A method for treating a subject for glaucoma, comprising administering a therapeutically effective amount of a deprenyl compound to a subject, wherein the deprenyl compound is represented by the structure:



in which

R<sub>1</sub> is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxy carbonyl, or aryloxy carbonyl;

A is a substituent independently selected for each occurrence from the group consisting of halogen, hydroxyl, alkyl, alkoxy, cyano, nitro, amino, carboxyl, -CF<sub>3</sub>, or azido;

n is 0 or an integer from 1 to 5;

and pharmaceutically acceptable salts thereof.

17.-20. **(Cancelled)**

21. **(Previously Presented)** The method of claim 14, wherein said therapeutically effective amount of a deprenyl compounds is administered with a pharmaceutically acceptable carrier.

22. **(Previously Presented)** The method of claim 21, wherein said pharmaceutically acceptable carrier has a pH of between about 3 and about 5.

23. **(Previously Presented)** The method of claim 22, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.

24.     **(Previously Presented)**       The method of claim 21, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.
25.     **(Previously Presented)**       The method of claim 14, wherein said subject is a human.
26.     **(Previously Presented)**       The method of claim 15, wherein said therapeutically effective amount of a deprenyl compounds is administered with a pharmaceutically acceptable carrier.
27.     **(Previously Presented)**       The method of claim 26, wherein said pharmaceutically acceptable carrier has a pH of between about 3 and about 5.
28.     **(Previously Presented)**       The method of claim 26, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.
29.     **(Previously Presented)**       The method of claim 28, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.
30.     **(Previously Presented)**       The method of claim 15, wherein said subject is a human.
31.     **(Previously Presented)**       The method of claim 16, wherein said therapeutically effective amount of a deprenyl compounds is administered with a pharmaceutically acceptable carrier.
32.     **(Previously Presented)**       The method of claim 31, wherein said pharmaceutically acceptable carrier has a pH of between about 3 and about 5.
33.     **(Previously Presented)**       The method of claim 31, wherein said pharmaceutically acceptable carrier is an alcohol or aqueous alcohol solution.

34.     **(Previously Presented)**     The method of claim 31, wherein said pharmaceutically acceptable carrier is suitable for ophthalmic administration.
35.     **(Previously Presented)**     The method of claim 16, wherein said subject is a human.